

-continued

Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	Trp	Tyr
	35						40					45			
Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	Glu	Glu
	50					55					60				
Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	Leu	His
65				70					75					80	
Gln	Asp	Trp	Leu	Asn	Gly	Lys	Glu	Tyr	Lys	Cys	Lys	Val	Ser	Asn	Lys
			85						90					95	
Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	Gly	Gln
			100					105					110		
Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	Glu	Leu
		115				120						125			
Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	Tyr	Pro
	130					135					140				
Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	Asn	Asn
145				150					155					160	
Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	Phe	Leu
			165					170						175	
Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	Asn	Val
		180					185						190		
Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	Thr	Gln
	195					200					205				
Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Lys							
	210					215									

1. An antigen binding construct comprising
 - a first antigen-binding polypeptide construct which monovalently and specifically binds a HER2 (human epidermal growth factor receptor 2) ECD2 (extracellular domain 2) antigen on a HER2-expressing cell;
 - a second antigen-binding polypeptide construct which monovalently and specifically binds a HER2 ECD4 (extracellular domain 4) antigen on a HER2-expressing cell;
 first and second linker polypeptides, wherein the first linker polypeptide is operably linked to the first antigen-binding polypeptide construct, and the second linker polypeptide is operably linked to the second antigen-binding polypeptide construct;
 and wherein the linker polypeptides are capable of forming a covalent linkage with each other.
- 2-35. (canceled)
36. The construct of claim 1, wherein the construct is conjugated to a drug.
37. The construct of claim 36, wherein the drug is maytansine (DM1).
38. (canceled)
39. A pharmaceutical composition comprising the construct of claim 1 and a pharmaceutical carrier.
40. The pharmaceutical composition of claim 39, the pharmaceutical carrier comprising a buffer, an antioxidant, a low molecular weight molecule, a drug, a protein, an amino acid, a carbohydrate, a lipid, a chelating agent, a stabilizer, or an excipient.
- 41-42. (canceled)

43. A method of treating a subject having a HER2 expressing (HER2+) tumor, comprising administering to the subject an effective amount of the construct of claim 1.
44. The method according to claim 43, wherein the result of the treatment is shrinking the tumor, inhibiting growth of the tumor, increasing time to progression of the tumor, prolonging disease-free survival of the subject, decreasing metastases, increasing the progression-free survival of the subject, or increasing overall survival of the subject.
45. The method according to claim 43, wherein the tumor is pancreatic cancer, head and neck cancer, gastric cancer, colorectal cancer, breast cancer, renal cancer, cervical cancer, ovarian cancer, endometrial cancer, uterine cancer, malignant melanoma, cancer of the pharynx, oral cancer or skin cancer.
- 46-59. (canceled)
60. A method of detecting or measuring HER2 in a sample comprising contacting the sample with the antigen binding construct according to claim 1 and detecting or measuring the bound complex.
61. A method of inhibiting, reducing or blocking HER2 signaling in a cell comprising administering an effective amount of the antigen binding construct according to claim 1 to the cell.
62. A method of killing or inhibiting the growth of a HER2-expressing tumor cell comprising contacting the cell with the antigen binding construct according to claim 1.
- 63-67. (canceled)
68. A method of producing the constructs according to claim 1 comprising culturing a host cell under conditions suitable for expressing the antigen binding construct